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NEWS 24 AUG 15 CAPLUS currency for Korean patents enhanced
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NEWS 28 SEP 25 CA/CAPLUS current-awareness alert options enhanced
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exemplified prophetic substances
NEWS 29 SEP 26 WPIDS, WPINDEX, and WPIX coverage of Chinese and
and Korean patents enhanced
NEWS 30 SEP 29 IFICLS enhanced with new super search field
NEWS 31 SEP 29 EMBASE and EMBAL enhanced with new search and
display fields
NEWS 32 SEP 30 CAS patent coverage enhanced to include exemplified
prophetic substances identified in new Japanese-
language patents

NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,
AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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FILE COVERS 1907 - 6 Oct 2008 VOL 149 ISS 15
FILE LAST UPDATED: 5 Oct 2008 (20081005/ED)

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=> s CB/ior-CEA.1

'TOR-CEA.1' IS NOT A VALID FIELD CODE
L1 0 CB/IOR-CEA.1

=> s (CB/ior-CEA.1)

'TOR-CEA.1' IS NOT A VALID FIELD CODE
L2 0 (CB/IOR-CEA.1)

=> s (CB and ior-CEA.1)
 14868 CB
 5020 CBS
 19503 CB
 (CB OR CBS)
 208 IOR
 7 IORS
 214 IOR
 (IOR OR IORS)
 7493 CEA
 216 CEAS
 7685 CEA
 (CEA OR CEAS)
 9833099 1
 2 IOR-CEA.1
 (IOR(W)CEA(W)1)
 L3 1 (CB AND IOR-CEA.1)

=> d L3 bib abs 1

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2003:892815 CAPLUS

DN 139:380012

TI Specific antibody fragments for the human carcinoembryonic antigen (CEA)

IN Gavilondo Cowley, Jorge Victor; Ayala Avila, Marta; Freyre Almeida, Freya de los Milagros; Acevedo Castro, Boris Ernesto; Bell Garcia, Hanssel; Roque Navarro, Lourdes Tatiana; Gonzalez Lopez, Luis Javier; Cremata Alvarez, Jose Alberto; Montesino Segui, Raquel

PA Centro de Ingenieria Genetica y Biotecnologia, Cuba

SO PCT Int. Appl., 49 pp.

CODEN: PIXXD2

DT Patent

LA Spanish

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2003093315	A2	20031113	WO 2003-CU5	20030428
WO 2003093315	A3	20040108		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,				

FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
 BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
 CA 2482411 A1 20031113 CA 2003-2482411 20030428
 AU 2003223831 A1 20031117 AU 2003-223831 20030428
 BR 2003004649 A 20040720 BR 2003-4649 20030428
 EP 1505076 A2 20050209 EP 2003-720119 20030428
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
 CN 1649901 A 20050803 CN 2003-809658 20030428
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 JP 2006500913 T 20060112 JP 2004-501454 20030428
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 ZA 2004008720 A 20060329 ZA 2004-8720 20041027
 MX 2004PA10695 A 20050217 MX 2004-PA10695 20041028
 US 20050158322 A1 20050721 US 2005-511794 20050317
 US 20070199078 A1 20070823 US 2007-731442 20070330
 PRAI CU 2002-86 A 20020429
 WO 2003-CU5 W 20030428
 US 2005-511794 A3 20050317

AB The invention relates to mono- and bivalent (diabody) single-chain Fv-type (scFv) antibody fragments which are obtained using recombinant DNA techniques from the carcinoembryonic anti-antigen (CEA) monoclonal antibody (McA) CB/ior-CEA.1. The aforementioned McA has a high affinity for the CEA and is used in the diagnosis and monitoring of colorectal tumors in humans. As with the original McA, diabody and monovalent scFv fragments exhibit high affinities for the human CEA and recognize an epitope that is dependent on carbohydrate conservation. The diabody and monovalent scFv fragments have affinity consts. for the CEA of $(5.0 \pm 0.4) \times 10^9$ L mol⁻¹ and $(2.8 \pm 0.3) \times 10^{10}$ L mol⁻¹ resp. The two aforementioned fragments do not display cross-reactivity with normal human tissues and cells, except for the normal colonic mucosa where the CEA is occasionally present. Said fragments can be produced through expression in recombinant micro-organisms from the cloning of nucleic acid sequences that code for variable regions obtained from the hybridoma that is produced by the CB/ior-CEA.1 McA. As with the original McA, the diabody and the monovalent scFv have a capacity for the in vivo identification in rats of human CEA-producing cells which grow forming tumors. The monovalent scFv and diabody do not possess Fc domains and the mol. sizes of said monovalent scFv and diabody are 5 and 2.5 times, resp., less than the rat McA. As a result, the aforementioned monovalent scFv and diabody can better penetrate tissues in vivo and are less immunogenic in humans.

=> s Tomoro and CEA

0 TOMORO

7493 CEA

216 CEAS

7685 CEA

(CEA OR CEAS)

L4 0 TOMORO AND CEA